## **Amendments to the Claims**

This listing of claims will replace all prior versions and listings of claims in the application.

### **Listing of Claims**

- 1. (Canceled)
- 2. (Currently Amended) A pharmaceutical composition according to claim-1 comprising a liposome formulated with at least one polypeptide, wherein the at least one polypeptide comprises an amino acid sequence at least 90% identical to the amino acid sequence set forth in SEQ ID NO: 2, wherein the liposome comprises a bacterial phospholipid, and wherein the composition is capable of inducing an immune response against Neisseria meningitidis.
- 3. (Currently Amended) The pharmaceutical composition according to elaim 4-claim 2, wherein the at least one polypeptide comprises an amino acid sequence at least 95% identical to the amino acid sequence set forth in SEQ ID NO: 2.
- 4. (Currently Amended) The pharmaceutical composition according to elaim 4claim 2, wherein the at least one polypeptide comprises the amino acid sequence set forth in SEQ ID NO: 2.
- 5. (Currently Amended) A pharmaceutical composition comprising a liposome formulated with a polypeptide comprising an immunogenic fragment comprising of at least 15-10 contiguous amino acids of SEQ ID NO: 2, wherein the liposome comprises a bacterial phospholipid, and wherein the composition is capable of inducing an immune response against *Neisseria meningitidis*.

# 6. (Canceled)

7. (Currently Amended) The pharmaceutical composition according to elaim +claim 2, wherein said at least one isolated polypeptide is selected from:

a polypeptide comprising the amino acid sequence set forth in SEQ ID NO:2 wherein the N-terminal methionine at residue 1 is deleted; and

a polypeptide comprising the amino acid sequence set forth in SEQ ID NO:2, wherein the secretory amino acid sequence is deleted.

### 8. -10. (Canceled)

- liposome formulated with a chimeric polypeptide that comprises two or more <u>immunogenic</u> fragments of a polypeptide, which polypeptide <u>comprises consists of</u> the amino acid sequence set forth in SEQ ID NO: 2, wherein each <u>immunogenic</u> fragment <u>is comprises</u> at least <u>15-10</u> contiguous amino acids of SEQ ID NO:2, and wherein the two or more <u>immunogenic</u> fragments are linked to form the chimeric polypeptide, <u>wherein the liposome is a bacterial phospholipid</u>, and wherein said composition is capable of inducing an immune response against *Neisseria* meningitidis.
- 12. (Currently Amended) The pharmaceutical composition according to elaim +claim 2, wherein the composition comprises at least two polypeptides wherein each of the at least two polypeptide polypeptides comprises an amino acid sequence at least 80%90% identical to the amino acid sequence set forth in SEQ ID NO: 2, and wherein the at least two polypeptides are linked to form a chimeric polypeptide.

# 13. (Canceled)

14. (Currently Amended) The pharmaceutical composition according to claim 13 any one of claims 2, 5, and 11, wherein said liposome comprises a the bacterial phospholipid is extracted from E. coli, N. meningitidis, or N. lactamica.

## 15. – 16. (Canceled)

- 17. (Currently Amended) The pharmaceutical composition according to elaim 13 claim 2, wherein said liposome further comprises at least one adjuvant selected from Lipid A, monophosphoryl lipid A (MPLA), a lipopolysaccharide, and a cytokine.
- 18. (Currently Amended) The pharmaceutical composition according to elaim 13claim 2, wherein said liposome <u>further</u> comprises 0 to 25% cholesterol.
- 19. (Currently Amended) The pharmaceutical composition according to any one of claims 1-5. 7, 11, and 122, 5, and 11, wherein said composition further comprises a pharmaceutically acceptable adjuvant.
- 20. (Currently Amended) A method for inducing an immune response against *N. meningitidis*; in a host, comprising administering to said host an immunogenic, effective amount of a-the pharmaceutical composition according to elaim lany one of claims 2, 5, and 11 to elicit an-the immune response.
- 21. (Currently Amended) A method for preventing or treating a N. meningitidis infection, said method comprising administering to a host in need thereof a prophylactic or therapeutic amount of a the pharmaceutical composition according to claim 1 any one of claims 2, 5, and 11.

### 22. (Canceled)

- 23. (Currently Amended) A method for the treatment or prophylaxis of meningitidis-meningitis and or meningococcemia caused by *Neisseria meningitidis* in a host, comprising administering to said host an effective amount of a the pharmaceutical composition according to elaim lany one of claims 2, 5, and 11.
- 24. (Currently Amended) A-The method according to claim 20, wherein said host is a mammal.
- 25. (Currently Amended) A-<u>The</u> method according to claim 24, wherein said host-mammal is a human.
- 26. (Currently Amended) A-The method according to claim 25, wherein said host human is an adult human.
- 27. (Currently Amended) A-The method according to claim 20 wherein the pharmaceutical composition is administered in unit dosage form of about 0.001 to 100 μg/kg (polypeptide weight/body weight) with an interval of about 1 to 6 weeks between immunizations.

### 28. –33. (Canceled)

- 34. (Currently Amended) The pharmaceutical composition according to any one of claims 12-5, 7, 11, and 12, wherein said polypeptide is capable of eliciting antibodies that are bactericidal.
- 35. (Currently Amended) The pharmaceutical composition according to any one of claims +2-5, 7, 11, and 12.

wherein the composition is capable of eliciting antibodies that bind to N. meningitidis of any one of serogroup A, B, and C.

- 36. (New) The pharmaceutical composition according to claim 5 wherein the immunogenic fragment comprises contiguous amino acids 108-125 of SEQ ID NO:2.
- 37. (New) The pharmaceutical composition of claim 36 wherein the composition further comprises a second immunogenic fragment, said second immunogenic fragment comprising contiguous amino acids 68-80 of SEQ ID NO:2.
- 38. (New) The pharmaceutical composition of claim 11 wherein one of the two or more immunogenic fragments comprises contiguous amino acids 108-125 of SEQ ID NO:2.
- 39. (New) The pharmaceutical composition of claim 11 wherein one of the two or more immunogenic fragments comprises contiguous amino acids 108-125 of SEQ ID NO:2 and one immunogenic fragment comprises contiguous amino acids 68-80 of SEQ ID NO:2.